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UTILITY PATENT APPLICATION TRANSMITTAL (Only for new nonprovisional applications under 37 CFR 1.53(b))	Attorney Docket No.	16904-738	Total Pages	34
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	Edward W. KNOWLTON, "Method and Apparatus for Controlled Contraction of Collagen Tissue"			
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APPLICATION ELEMENTS See MPEP chapter 600 concerning utility patent application contents.	ADDRESS TO: Assistant Commissioner for Patents Box Patent Application Washington, DC 20231
1. <input type="checkbox"/> Fee Transmittal Form (Submit an original, and a duplicate for fee processing)	6. <input type="checkbox"/> Microfiche Computer Program (Appendix)
2. <input checked="" type="checkbox"/> Specification [Total Pages <u>27</u>] (preferred arrangement set forth below) <ul style="list-style-type: none">- Descriptive title of the Invention- Cross References to Related Applications- Statement Regarding Fed sponsored R&D- Reference to Microfiche Appendix- Background of the Invention- Brief Summary of the Invention- Brief Description of the Drawings- Detailed Description- Claim(s)- Abstract of the Disclosure	7. Nucleotide and/or Amino Acid Sequence Submission (if applicable, all necessary) <ul style="list-style-type: none">a. <input type="checkbox"/> Computer Readable Copyb. <input type="checkbox"/> Paper Copy (identical to computer copy)c. <input type="checkbox"/> Statement verifying identify of above copies
3. <input checked="" type="checkbox"/> Drawing(s) (37 CFR 1.152) [Total Sheets <u>6</u>]	8. <input type="checkbox"/> Assignment Papers (cover sheet & documents(s))
4. <input type="checkbox"/> Oath or Declaration [Total Pages <u> </u>] <ul style="list-style-type: none">a. <input type="checkbox"/> Newly executed (original or copy)b. <input type="checkbox"/> Copy from a prior application (37 CFR 1.63(d)) (for continuation/divisional with Box 17 completed) [Note Box 5 below]<ul style="list-style-type: none">i. <input type="checkbox"/> DELETION OF INVENTOR(S) Signed statement attached deleting inventor(s) named in the prior application, see 37 CFR 1.63(d)(2) and 1.33(b).	9. <input type="checkbox"/> 37 CFR 3.73(b) Statement [] Power of Attorney (when there is an assignee)
5. <input type="checkbox"/> Incorporation By Reference (useable if Box 4b is checked) The entire disclosure of the prior application, from which a copy of the oath or declaration is supplied under Box 4b, is considered as being part of the disclosure of the accompanying application and is hereby incorporated by reference therein.	10. <input type="checkbox"/> English Translation Document (if applicable)
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17. If a **CONTINUING APPLICATION**, check appropriate box and supply the requisite information:☒ Continuation [] Divisional [] Continuation-in-part (CIP) of prior application No. 08 / 914,681**16. CORRESPONDING ADDRESS**

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**UNITED STATES
PATENT APPLICATION**

entitled

**METHOD AND APPARATUS FOR CONTROLLED
CONTRACTION OF COLLAGEN TISSUE**

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Status: Small Entity

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**METHOD AND APPARATUS FOR CONTROLLED
CONTRACTION OF COLLAGEN TISSUE**

Cross-Reference to Related Cases

5 The present application is a continuation of U.S. Patent Application
Serial No. 08/914,681, filed August 17, 1997, which is a continuation of
U.S. Patent Application Serial No. 08/435,822, filed May 5, 1995, entitled
"Method and Apparatus for Controlled Contraction of Collagen Tissue", and
10 is related to U.S. Patent Application Serial No. 08/435,544 filed May 5,
1995, entitled "Method and Apparatus for Controlled Contraction of
Collagen Tissue", all applications having the same named inventor Edward
W. Knowlton, and all applications being incorporated herein by reference.

BACKGROUND OF THE INVENTION

Field of the Invention

15 This invention relates generally to a method and apparatus for
shrinking skin tissue without damaging the melanocytes and other epithelial
cells, and more particularly to a method and apparatus that creates a reverse
thermal gradient in order to achieve skin tightening through partial
denaturation of collagen without ablation of the collagen and without
20 affecting the melanocytes and other epithelial cells.

Description of Related Art

 The skin is the one organ of the body that is readily available for
inspection by the eyes and fingers of every living person. It is subjected to

considerable abuse such as exposure to extreme environmental of cold, heat, wind, and sun.

The surface layer of the skin is called the epidermis. It is the barrier that prevents most substances from entering the body from outside while preventing body fluids from entering equilibrium with the environment. The basilar layer of the epidermis includes the melanocytes and other epithelial cells.

The melanocytes are small cells with a small, dark staining nucleus and a clear cytoplasm. Melanin is packaged in distinctive granules is produced by these cells and transferred then air dendritic processes to adjacent keratinocytes. The purpose of melanin is to protect the skin from the ravages of ultraviolet radiation. Beneath the epidermis is the papillary dermis and reticular dermis. Collagen tissue is found in the dermal and the sub dermal tissues.

There has been a large market for tightening the skin in order to reduce aging effects and effects created by exposing the skin to extreme environmental forces. To date there are two primary methods for tightening skin. The first is surgical skin excision. The second is chemical burn. When skin is surgically excised it leaves large scares. This is generally not a satisfactory solution for many skin tightening applications. With chemical peel treatments the process is painful, there is damage to the melanocytes and other epithelial cells, the patient maybe have spotted pigmentation, or with most of the melanocytes and other epithelial cells destroyed the patient can have a predominately white complexion. In the chemical peel method a thermal gradient is created which is hotter at the surface of the epidermis and cooler at the sub dermal layers. With the creation of this type of thermal

gradient there is a great likelihood of modification or destruction of the melanocytes and other epithelial cells, resulting in blotchiness or an inability to tan in the future.

Collagen molecules are produced by fibroblasts which synthesize three polypeptide chains that wrap around one another in a triple helix. Each of the chains is approximately 1000 amino acid units in length, with glycine recurring regularly every third unit and hydroxyproline and proline recurring very frequently. Cross-linking occurs between the side, not the ends, of collagen molecules and is coupled with the amino acid composition to give collagen its great strength. Collagen tissue shrinkage takes place in a direction parallel to an axis of collagen fibers.

The phenomenon of thermal shrinkage of collagen begins with a denaturation of the triple helix of the collagen molecule. Partial denaturation of collagen tissue results in a shrinkage of the collagen and provides a "tightening" effect on the overlaying skin. To date there have been no devices or methods for contracting the underlying collagen tissue through partial denaturation without damaging the melanocytes and other epithelial cells in the epidermis.

Adipose tissue, more commonly known as fat, is formed of cells containing stored lipid. Adipose tissue is often subdivided into small lobules by connective collagen tissue serving as the fibrous septae.

Adipose tissue is widely distributed in the subcutaneous tissue but exhibits regional differences in amount partially because of age and sex. Excess adipose tissue can be physically undesirable from both health and cosmetic perspective. A current method for the removal of fatty tissue is the extraction of adipose tissue by liposuction. This is a purely mechanical

method with undesirable side effects due to the invasive nature of the process.

Another method of removing fatty tissue is disclosed in U. S. Patent No. 5,143,063 ("the '063"). The method of the '063 patent targets adipose tissue absorbs sufficient energy resulting in cell destruction and death. The method of the '063 patent fails to minimize damage to the melanocyte in the epidermis. Thus, with the method of the '063 patent can create unwanted blotchiness or changes in the melanocytes and other epithelial cells.

There exists the need for skin tightening without damaging the melanocytes and other epithelial cells, or without surgical intervention. There is a further need for non- surgically removing adipose tissue without damaging the melanocytes and other epithelial cells.

SUMMARY OF THE INVENTION

It is an object of the present invention to provide a method and apparatus for tightening skin without substantially damaging the melanocytes and other epithelial cells.

Another object of the present invention is to provide a method and apparatus for tightening skin with the use of a thermal energy source that does not substantially effect the melanocytes and other epithelial cells.

Yet another object of the invention is to provide an inverse thermal gradient from the epidermis to the underlying collagen tissue.

A further object of the invention is to provide a method and apparatus for creating a reverse thermal gradient from the epidermis through a subcutaneous layer of soft tissue.

Still a further object of the present invention is to provide a method and apparatus for tightening skin through the use of an RF energy source and an electrolytic solution which transfer RF energy from RF electrodes to the epidermis and underlying layers.

5 Yet another object of the present invention is to provide a method and apparatus for creating a reverse thermal gradient that utilizes one or more RF electrodes, an electrolytic solution to transfer RF energy from the RF electrodes to the epidermis and underlying layers, and wherein the apparatus includes a lumen adapted to receive a cooling fluid.

10 These and other objects of the invention are provided in an apparatus for applying radiant energy through the skin to underlying collagen tissue without substantially modifying melanocytes and other epithelial cells in the epidermis. The apparatus includes a porous membrane adapted to receive an electrolytic solution and become inflated to substantially conform a
15 contacting exterior surface of the membrane to a skin layer. The membrane includes a cooling lumen for receiving a cooling fluid. One or more thermal electrodes are positioned in the membrane and transfer thermal energy to the electrolytic solution. The electrolytic solution and cooling fluid create a reverse thermal gradient from the skins surface to the underlying collagen
20 tissue. A thermal power source is coupled to the thermal electrodes, and a source of electrolytic solution is coupled to the membrane.

25 A method for tightening skin comprises providing a membrane in a thermal energy source. A reverse thermal gradient is created which cools the surface of the skin while heating underlying collagen containing layers. The skin and underlying collagen containing tissue are then heated without substantially modifying the melanocytes and other epithelial cells in the

epidermis. The result is a contraction of collagen tissue and a tightening of the skin.

Radiant energy is applied to a variety of different skin layers including the papillary dermis layer, the reticular dermis layer, and even to a subcutaneous layer and to underlying soft tissue. One suitable energy source is one or more RF electrodes. Electrolytic solution transfers RF energy from the RF electrodes to the underlying collagen tissue. The cooling fluid can create a reverse thermal gradient at the epidermis to underlying desired layers of about 30 degrees to about 80 degrees C. The apparatus can further include one or more thermal sensors positioned on the contacting exterior surface of the membrane, as well as one or more impedance monitors. Further, the apparatus can include a feedback device which is responsive to a detected characteristic of the skin or subcutaneous layer in order to provide a control delivery of RF energy from the RF electrodes. A variety of detected characteristics can be monitored including impedance measurements of the skin and temperature profiles of the skin.

The feedback device can also include a controller as well as a multiplexer.

The creation of the reverse thermal gradient provides for the controlled contraction of collagen tissue, e.g., partial denaturization of the collagen molecules that results in a shrinkage of the collagen tissue, which then extends to a shrinkage of the skin. Creation of the reverse thermal gradient is different from other methods of collagen contraction which typically employ a thermal gradient that has a higher temperature at the surface and decreases with the depth of penetration.

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The apparatus of the present invention that creates the desired contour effect by the modification of collagen tissue surrounding loculations of fat can include a focussing element to help direct the energy past the epidermis and into the fat containing layers. The apparatus can further include a coupling device that couples the focussing element to the membrane, as well as a bracket which is positioned around the periphery of the membrane and also supports the focussing elements. Instead of killing cells, thermal lipolysis is utilized with a cooling blanket on the skin so that thermal destruction fat cells, and the melanocytes and other epithelial cells are minimally damaged.

A variety of thermal energy sources can be employed. Suitable energy sources include but are not limited to RF, microwave, ultrasound and the like. In one embodiment, the preferred energy source is RF.

BRIEF DESCRIPTION OF THE FIGURES

Fig. 1 is a perspective view of an apparatus for applying radiant energy through the skin in order to cause a partial denaturization of collagen tissue, resulting in a tightening of the skin.

Figure 2 is a cross-sectional view of the skin and underlying tissue.

Figure 3 is a schematic representation of the collagen network.

Fig. 4 is a schematic diagram of an apparatus for applying radiant energy to underlying subcutaneous layers or deeper soft tissue layers to create a desired contour effect by partially denaturing collagen tissue, and without substantially modifying melanocytes and other epithelial cells in the epidermis.

Figure 5 is a block diagram of an RF system which can be utilized with the present invention.

Figure 6 is a block diagram of processing circuit of one embodiment of the invention.

DETAILED DESCRIPTION

The present invention provides an apparatus for applying radiant energy through the skin to underlying collagen tissue without substantially modifying the melanocytes and other epithelial cells found in the epidermis.

The present invention also provides an apparatus for applying radiant energy through the skin to underlying subcutaneous or deeper soft tissue layers that include loculations of fat with fibrous septae made of collagen tissue. Application of the radiant energy creates an desired contour effect of the loculations of fat without substantially modifying the melanocytes and other epithelial cells in the epidermis. Further, non invasive methods are providing for skin tightening and tightening the fibrous septae around the loculations of fat to create the desired contour effect.

The methods of the present invention do not provide for total necrosis of cells. Instead, with the method and apparatus creating the desired contour effect, the loculations of fat with the fibrous septae made of collagen tissue use a reverse thermal gradient applied to the underlying collagen tissue layers resulting in a partial denaturization of the collagen permitting it to become tightened. This is achieved without killing all of the fat cells within the loculations.

Various types of radiant energy can be utilized with the present invention. Radiant energy may be any kind that can cause cell heating or

physical destruction by being applied to collagen tissue. Examples of suitable radiant energy source include, but are not limited to RF, microwave, ultrasound, and the like.

Referring now to Fig. 1, an apparatus 10 applies radiant energy through a skin layer 12, such as the epidermis, and to the underlying collagen tissue 14 without substantially modifying melanocytes and other epithelial cells 16 found in the lower layer of epidermis layer 12.

A porous membrane 18 is adapted to receive an electrolytic solution 20. Porous membrane 18 becomes inflated to substantially conform a contacting exterior surface 22 of porous membrane 18 which is in close thermal contact with epidermis 12. Porous membrane 18 includes a cooling lumen 24 for receiving a cooling fluid that imparts a cooling effect on epidermis layer 12.

One or more thermal electrodes 26 are positioned at various places in porous membrane 18. In one embodiment, thermal electrodes 26 are positioned on a side that is substantially opposing to contacting exterior surface 22. In other embodiments, thermal electrodes 26 are placed closer to cooling lumen 24. In embodiment particularly suitable for the hips, porous membrane is about 20 cm by 30 cm, with an oval shape.

A thermal power source 28 is coupled to thermal electrodes 26 and a source of electrolytic solution 30 is coupled to porous membrane 18.

With referenced now to Fig. 2, radiant energy can be applied through epidermis layer 12, to papillary dermis layer 32, to reticular dermis layer 34, to subcutaneous layer 35, as well as to underlying soft tissue 36. The extend of collage in the various layers is < 5% in the epidermis, ~ 50% in the dermis, ~ 20 % in the subcutaneous, ~ 5% in the muscle with overlying

fascia. Shrinking of collagen tissue takes place in a direction parallel to the axis of the collagen fibers. Thermal shrinkage of collagen begins with the denaturization of the triple helix structure of the collagen fibers. This occurs when thermal energy is applied to the collagen tissue causing the hydrolysis of heat labial cross links of the collagen network.

Fig. 3 is a schematic representation of a collagen network behavior under the influence of heat. The thickened lines represent the chains originally bound by covalent cross links. The arrows indicate tensions exerted on the collagen chains by the effect of heat. More particularly, Fig. 3 illustrates (i). native collagen network 40, (ii). collagen 42 under isometric conditions, (iii). collagen network without any restraint, (iv). collagen network 46 under isometric tension as long as the nodes are stable, and (v). collagen network 48 under isometric tension after some cross links have been cleaved.

In one embodiment of the present invention, thermal electrodes 26 are RF electrodes which can be a single electrode, or a plurality which can form a segmented flexible circuit. Thermal power source 28 is then an RF generator. Electrolytic solution 20 is introduced into porous membrane 18 and passes by RF electrodes 26. Electrolytic solution 20 transfers RF power from RF electrodes 28 to the desired underlying collagen tissue to achieve partial denaturization of the collagen molecule.

Generally, RF electrodes 26 can be monopolar or bipolar. In the monopolar mode, RF current flows through body tissue from a return electrode which can be in a form of a conductive pad applied to the patients outer skin. Maximum heating occurs where the current density is the greatest.

During a treatment phase, the denaturization of collagen molecules can be conducted under feedback control. Treatment can occur without the attention of medical supervision. Feedback is accomplished by (i). visualization, (ii). impedance, (iii). ultrasound, or (iv). temperature measurement. Optionally included and preferably positioned on contacting exterior surface 22 can be one or more thermal sensors 52, as well as one or more impedance monitors 54. Thermal sensors 52 permit accurate determination of the surface temperature of epidermis layer 12.

Electrolytic solution 20 can be preheated to a selected temperature and modified as necessary. This reduces the amount of time needed to effect at satisfactory denaturization of collagen molecules and subsequent skin tightening.

Porous membrane 18 can be made of a material that is an insulator. For purposes of this disclosure, an insulator is a barrier to thermal or electrical energy flow. Porous membrane 18 can be made of a material which permits controlled delivery of electrolytic solution 20 to epidermis layer 12. Porous membrane 18 can be made of a variety of materials including, but not limited to knitted polyester, continuous filament polyester, polyester-cellulose, rayon, polyamide, polyurethane, polyethylene and the like. Suitable commercial products include, (i). Opcell available from Centinal Products Corp., Hyannis, Mass, and (ii). UltraSorb, HC 4201 or HT 4644 MD from Wilshire Contamination Control, Carlsbad, California. Pockets or zones 56 can be formed around RF electrodes 26. Each pocket 56 has a lower porosity for the flow of electrolytic solution 20 than all other sections of porous membrane 18. Differences in porosity can be achieved with different types of materials which form porous membrane 18.

Electrolytic solution 20 is retained in pockets 56 longer than in non-pocket sections of porous membrane 18, and there is a greater transfer of RF energy to electrolytic solution 20, creating a larger electrode. The larger electrode produces RF and thermal energy to create a larger electrode effect.

5 However, this does not effect the creation of the reverse thermal gradient. RF energy is still transferred through porous membrane 18 passing in the vicinity of cooling lumen 24, in order to create a lower temperature at epidermis layer 12 and the temperature increases as deeper layers are reached.

10 Referring now to Fig. 4, an apparatus 58 for creating a desired contour effect of underlying subcutaneous layers or deeper soft tissue layers which include loculations of fat with fibrous septae made of collagen tissue is illustrated. The apparatus 58 of Fig 4, includes a porous membrane 18, electrolytic solution 20, a contacting exterior surface 22, a cooling lumen, 15 thermal electrodes 26, a thermal power source 28, an electrolytic solution source 30, one or more thermal sensors 52, as well as one or more impedance monitors 54. Apparatus 58 also includes a focussing element 60 which focuses thermal energy from electrolytic solution 20 to the underlying collagen tissue. Focussing element 60 and electrolytic solution 20 create a 20 reverse thermal gradient from epidermis layer 12 to the underlying collagen tissue 14. Focussing element 62 can be, in the case of ultrasonic energy, a lens having a flat planer surface on the radiation wave incident side and a concave exit face, see *Ultrasonics Theory and Application*, by G.L. Goberman, Heart Publishing Co., New York (1959), at section 2.6. The use 25 of such a focussing lens for ultrasonic energy with a planer wave receiving face and concave exit face is also described in the article "Deep Local

Hypothermia for Cancer Therapy: Extreme Electromagnetic and Ultrasound
Technics," A.Y. Cheung and A. Neyzari, *Cancer Research*, Vol. 44,
pp.4736-4744, October 1984.

Radio frequencies can be the thermal energy source, and various
5 localizing technique, well known in the art, can be utilized. In one
embodiment, radio frequency energy is supplied by capacitive coupling
directly to epidermis layer 12 for areas close to the dermal tissue. Radio
frequency induction focussing can be achieved with the use of plural
focussing coils which are adaptive at the zone of interest and are elsewhere
10 subtractive. Alternatively, radio frequency energy may be focused by having
a multiple beam phased array. For concave focussing see, "Tumor reduction
by radio frequency therapy response", H.H. Lavien et al., *JAMA*, Vol. 233,
at 2198-2200.

Alternative radio frequency focussing methods are disclosed in
15 "Equipment for Local Hypothermia Therapy of Cancer", C.F. Babbs et al.,
Medical Instrumentation, Vol. 16, No. 5, Sept-Oct 1982, pp.245-248.

It will be appreciated that focussing element 60 can be a convergent
lens. Further, focussing element 60 can be positioned in porous membrane
18, and at the exterior 16 between epidermis layer 12 and porous membrane
20 18. Further, a coupling device 62 can be included which couples focussing
element 60 with porous membrane 18. In one embodiment, coupling device
62 is a bracket which is positioned around a periphery of porous membrane
18, and supports focussing element 50 in relation to porous membrane 18.

In the method for tightening skin, porous membrane 18 and thermal
25 energy source 26 are provided. A reverse thermal gradient is created which
cools a surface of epidermis layer 12 while heating underlying collagen

containing layers. Epidermis layer 12 as well as underlying collagen containing tissue are heated, without substantially effecting the melanocytes and other epithelial cells in epidermis layer 12, resulting in a denaturization of collagen molecules, causing a contraction of the collagen tissue and a tightening of the skin. This method can be applied numerous times. In many instances, it may be desirable to tighten the skin to a certain level and then in subsequent treatments the skin is tightened further. There are may be four fine treatments to fine tune the contour effects with greater precision. In this method, collagen containing tissue is partial denatured and fat cell destruction is minimized. This is achieved by partially denaturing by cleaving heat labial cross links of the collagen molecules.

The reverse thermal gradient provides a variation in temperature throughout the various tissue layers. For example, in one embodiment, the reverse thermal gradient is a temperature range from about 30 degrees to about 80 degrees C, initiating from epidermis layer 12 to collagen containing tissue layers. The reverse thermal gradient can have a temperature range of about 30 to 75 degrees C, and additionally from about 30 to 70 degrees C.

In another embodiment, a method for liposculpturing an area of the body where there is an underlying area comprised of a loculation of fat that has collagen tissue as a fibrous septae also includes creating a reverse thermal gradient from epidermis layer 12 to the desired underlying loculation of fat layer. Sufficient thermal energy is supplied through epidermis layer 12, without damaging or substantially modifying the melanocytes and other epithelial cells, through other skin layers and is focused on the collagen tissue of the fibrous septae. Thermal energy partially denatures the collagen tissue with a minimal destruction of fat cells. Again, this is achieved by partially

denaturizing, e.g., by cleaving, heat labial cross links of collagen molecules. The reverse thermal gradient produces a net mobilization of intra-cellular fat with diminished destruction of fat cells.

For purposes of illustration, without the scope of the invention let it be assumed that power source 28 is an RF power source, an RF power source 28, feeds energy to an RF power generator 64 and then to RF electrodes 26. A multiplexer 66 measures current, voltage and temperature, at the numerous thermal sensors associated with to each RF electrode 26. RF electrodes 26 can be individually measured. Multiplexer 66 is driven by a controller 68 which can be a digital or analog controller, or a computer with software. When controller 68 is a computer it can include a CPU coupled through a system bus. On the system can be a keyboard, disk drive, or other non volatile memory systems, a display, and other peripherals, as are well known in the art. Also coupled to the bus are a program memory and a data memory.

An operator interface 70 includes operator controls 72 and a display 74. Controller 68 can be coupled to different types of imaging systems including ultrasonic, thermal sensors 52, and impedance monitors 54.

Current and voltage are used to calculate impedance. A diagnostic phase can be initially run to determine the level of treatment activity. This can be done through ultrasound as well as other means. Diagnostics can be performed both before and after treatment.

Thermal sensors 52, and thermal sensors 76 contained within RF generator 64 measure voltage and current that is delivered to the desired treatment site. The output for these sensors is used by controller 68 to control the delivery of RF power. Controller 68 can also control

temperature and power. An operator set level of power and/or temperature may be determined and this will not be exceeded. Controller 68 maintains the set level under changing conditions. The amount of RF energy delivered controls the amount of power. A profile of power delivered can be
5 incorporated in controller 68, as well as a preset amount of energy to be delivered. Feedback can be the measurement of impedance, temperature, or other indicators and occurs either at control 68 or at RF generator 64, if it incorporates a controller. For impedance measurement, this can be achieved by supplying a small amount of non therapeutic RF energy. Voltage and
10 current are then measured to confirm electrical contact.

Circuitry, software and feedback to controller 68 result in full process control and are used to change, (i). power, (ii). the duty cycle, (iii). monopolar or bipolar energy delivery, (iv). electrolytic solution 20 delivery, flow rate and pressure and (v). can determine when the process is completed
15 through time, temperature and/or impedance. These process variables can be controlled and varied based upon tissue temperature monitored at multiple sites on contacting exterior surface 22 as well as monitoring impedance to current flow at each RF electrode 26, indicating changes in current carrying capability of the tissue during the process. Further, controller 68 can provide
20 multiplexing, monitor circuit continuity, and determine which RF electrode 26 is activated.

A block diagram of one embodiment of suitable processing circuitry is shown in Fig. 6. Thermal sensors 52 can be thermistors which have a resistance that varies with temperature. Analog amplifier 78 can be a
25 conventional differential amplifier circuit for use with thermistors and transducers. The output of analog amplifier is sequentially connected by an

analog multiplexer 80 to the input of an analog digital converter 82. The output of amplifier 78 is a voltage which represents the respective sensed temperatures. The digitized amplifier output voltages are supplied by analog to digital converter 82 to a microprocessor 84. Microprocessor 84
5 calculates the temperature or impedance of the tissue. Microprocessor 84 can be a type 6800. However, it will be appreciated that any suitable microprocessor or general purpose digital or analog computer can be used to calculate impedance or temperature.

Microprocessor 84 sequentially receives and stores digital
10 representations of impedance and temperature. Each digital value received by microprocessor 84 corresponds to different temperatures and impedances.

Calculated temperature and impedance values can be indicated on display 74. Alternatively, or in addition to the numerical indication of temperature or impedance, calculated impedance or temperature values can
15 be compared by microprocessor 84 with temperature and impedance limits. When the values exceed predetermined temperature or impedance values a warning can be given on display 74 and additionally, the delivery of RF energy to its respective electrode can be decreased or multiplexed to another electrode. A control signal from microprocessor 84 can reduce the power
20 level by RF generator 64, or de-energize the power delivered to any particular electrode. Controller 68 receives and stores the digital values which represent temperatures and impedances sent. Calculated surface temperatures and impedances can be forwarded by controller 68 to display 74. If desired, the calculated surface temperature of epidermis layer 12 is
25 compared with a temperature limit and a warning signal can be sent to

display 74. Similarly, a control signal can be sent to RF power source 26 when temperature or impedance values exceed a predetermined level.

The foregoing description of a preferred embodiment of the invention has been presented for purposes of illustration and description. It is not intended to be exhaustive or to limit the invention to the precise forms disclosed. Obviously, many modifications and variations will be apparent to practitioners skilled in this art. It is intended that the scope of the invention be defined by the following claims and their equivalents.

CLAIMS

1 1. A skin treatment apparatus, comprising:
2 a fluid delivery member with a tissue interface surface that remains
3 conformable to a skin surface as the tissue interface surface is applied to a
4 surface of the skin; and
5 a thermal energy delivery device coupled to the membrane in a
6 position to transfer thermal energy to an electrolytic media that passes
7 through the fluid delivery member.

1 2. The apparatus of claim 1, wherein the thermal energy delivery
2 device is positioned in an interior of the fluid delivery member.

1 3. The apparatus of claim 1, wherein the thermal energy delivery
2 device is positioned at an exterior surface of the fluid delivery member.

1 4. The apparatus of claim 1, wherein the thermal energy delivery
2 device is positioned at the tissue interface surface.

1 5. The apparatus of claim 1, wherein the thermal energy delivery
2 device is at least one RF electrode.

1 6. The apparatus of claim 1, further comprising:
2 a fluid passage lumen coupled to the fluid delivery member.

1 7. The apparatus of claim 1, wherein the tissue interface surface
2 of the fluid delivery member has a porous surface.

1 8. The apparatus of claim 1, wherein at least a portion of the
2 fluid delivery member is a membrane.

1 9. The apparatus of claim 1, further comprising:
2 a cooling fluid lumen coupled to the fluid delivery member.

1 10. The apparatus of claim 1, further comprising:
2 a sensor coupled to the fluid delivery member.

1 11. The apparatus of claim 10, wherein the sensor is positioned at
2 the tissue interface surface of the fluid delivery device.

1 12. The apparatus of claim 10, further comprising:
2 a feedback device coupled to the energy delivery device and the
3 sensor, the feedback device being responsive to a detected characteristic of a
4 tissue site and provide a controlled delivery of thermal energy.

1 13. An apparatus for treating a skin surface, comprising:
2 a fluid receiving member;
3 a thermal energy delivery device coupled to the fluid receiving
4 member;
5 an electrolytic media positionable in the fluid receiving member, the
6 thermal energy delivery device being positioned in the fluid receiving

7 member to transfer thermal energy to the electrolytic media, wherein a
8 selected collagen containing tissue site under the skin surface receives the
9 thermal energy and creates a tightening of the skin surface.

1 14. The apparatus of claim 13, wherein the fluid receiving
2 member includes a tissue interface surface.

1 15. The apparatus of claim 14, wherein at least a portion of the
2 tissue interface is porous.

1 16. The apparatus of claim 13, wherein at least a portion of the
2 fluid receiving member is a membrane.

1 17. The apparatus of claim 13, wherein the fluid receiving
2 member includes a conformable tissue interface surface.

1 18. The apparatus of claim 17, wherein the tissue interface
2 surface is conformable to the skin surface.

1 19. The apparatus of claim 17, wherein the tissue interface
2 surface remains conformable to a skin surface as the tissue interface surface
3 is applied to a surface of the skin.

1 20. The apparatus of claim 13, wherein the thermal energy
2 delivery device is positioned in an interior of the fluid delivery member.

1 21. The apparatus of claim 13, wherein the thermal energy
2 delivery device is positioned at an exterior surface of the fluid delivery
3 member.

1 22. The apparatus of claim 13, wherein the thermal energy
2 delivery device is at least one RF electrode.

1 23. The apparatus of claim 1, further comprising:
2 a fluid passage lumen coupled to the fluid delivery member.

1 24. The apparatus of claim 1, wherein the tissue interface surface
2 of the fluid delivery member has a porous surface.

1 25. The apparatus of claim 1, wherein at least a portion of the
2 fluid delivery member is a membrane.

1 26. The apparatus of claim 1, further comprising:
2 a cooling fluid lumen coupled to the fluid delivery member.

1 27. The apparatus of claim 1, further comprising:
2 a sensor coupled to the fluid delivery member.

1 28. The apparatus of claim 27, wherein the sensor is positioned of
2 the tissue interface surface of the fluid delivery device.

1 29. The apparatus of claim 27, further comprising:

2 a feedback device coupled to the energy delivery device and the
3 sensor, the feedback device being responsive to a detected characteristic of a
4 tissue site and provide a controlled delivery of thermal energy.

1 30. A method for tightening an external surface of a skin with an
2 underlying collagen containing tissue, comprising:

3 providing a thermal energy delivery device with an energy delivery
4 surface;

5 positioning the thermal energy delivery surface on the external
6 surface of the skin;

7 creating a reverse thermal gradient through the surface of the skin to
8 sufficiently heat an underlying collagen containing tissue, wherein a
9 temperature of the external surface of the skin is lower than a temperature of
10 the underlying collagen containing tissue;

11 detecting a temperature of the external surface of the skin;

12 heating the underlying collagen containing tissue in response to a
13 detected temperature of the external surface of the skin; and

14 tightening at least a portion of the external surface of the skin.

1 31. A method for tightening skin, comprising:

2 providing a thermal energy delivery device;

3 positioning an energy delivery surface of the thermal energy delivery
4 device on an external surface of the skin;

5 creating a reverse thermal gradient through a surface of the skin
6 while heating underlying collagen containing tissue, wherein a temperature of

7 the external skin surface is lower than a temperature of the underlying
8 collagen containing tissue;
9 heating the underlying collagen containing tissue without creating a
10 necrosis of living cells in the epidermis;
11 contracting at least a portion of the collagen containing tissue; and
12 tightening at least a portion of the surface of the skin.

1 32. A method for tightening skin, comprising:
2 providing a thermal energy delivery device;
3 positioning an energy delivery surface of the thermal energy delivery
4 device on an external surface of the skin;
5 creating a reverse thermal gradient through a surface of the skin
6 while heating underlying collagen containing tissue, wherein a temperature of
7 the external skin surface is lower than a temperature of the underlying
8 collagen containing tissue;
9 controlling a delivery of a sufficient amount of thermal energy
10 through an epidermis of the external surface of the skin to reconfigure at
11 least a portion of an underlying collagen containing tissue without
12 substantially creating cell necrosis in the epidermis, wherein at least a portion
13 of the surface of the skin is tightened.

1 33. A method for tightening skin, comprising:
2 providing a thermal energy delivery device;
3 positioning an energy delivery surface of the thermal energy delivery
4 device on an external surface of the skin;

5 heating through a surface of the skin the collagen containing tissue
6 underlying the surface of the skin, wherein a temperature of the external skin
7 surface is lower than a temperature of the underlying collagen containing
8 tissue;

9 controlling a delivery of a sufficient amount of thermal energy
10 through an epidermis of the surface of the skin to reconfigure at least a
11 portion of an underlying collagen containing tissue without substantially
12 creating cell necrosis in the collagen containing tissue, wherein at least a
13 portion of the surface of the skin is tightened.

1 34. An apparatus for applying energy through a skin epidermis
2 surface of an underlying subcutaneous layer or deeper soft tissue layers that
3 includes collagen containing tissue, comprising:

4 a membrane that conforms a contacting exterior surface of the
5 membrane to the skin epidermis surface;

6 one or more electrodes positioned in the membrane configured to be
7 coupled to an energy source;

8 an electrolytic media positioned in the membrane a coupled to the
9 electrodes to receive energy from the electrodes and transfer energy from the
10 electrodes to the skin epidermis surface;

11 a focussing element that creates a reverse thermal gradient from the
12 skin epidermis surface to the collagen containing tissue.

1 35. An apparatus for applying energy through a skin epidermis
2 surface to an underlying collagen containing tissue; comprising:

3 an energy delivery device means;

4 electrolytic media energy delivery means coupled to the energy delivery
5 device means to receive electrolytic energy and transfer the electrolytic
6 energy from the energy delivery device means to the skin surface;
7 the electrolytic media means delivering energy to the skin surface and
8 the energy passing through the skin surface to the underlying collagen
9 containing tissue to contract at least a portion of the collagen containing
10 tissue without substantial cell necrosis and creating a tightening of the skin
11 surface; and
12 a membrane housing means housing at least a portion of the energy
13 delivery means, the membrane means including a membrane skin surface
14 interface means.

ABSTRACT

METHOD AND APPARATUS FOR CONTROLLED CONTRACTION OF COLLAGEN TISSUE

5 A method and apparatus for tightening a surface of a skin with an
underlying collagen containing tissue applies radiant energy through the skin
to underlying collagen tissue without substantially modifying melanocytes
and other epithelial cells in the epidermis. A porous membrane is adapted to
receive an electrolytic solution and become inflated to substantially conform
a contacting exterior surface of the membrane to a skin layer. The
10 membrane includes a cooling lumen for receiving cooling fluid. One or more
thermal electrodes positioned in the membrane and transfers thermal energy
to the electrolytic solution. The electrolytic solution and cooling fluid
creates a reverse thermal gradient from the skin surface to the underlying
collagen tissue. A thermal power source is coupled to the thermal
electrodes, and a source of electrolytic solution is coupled to the membrane.

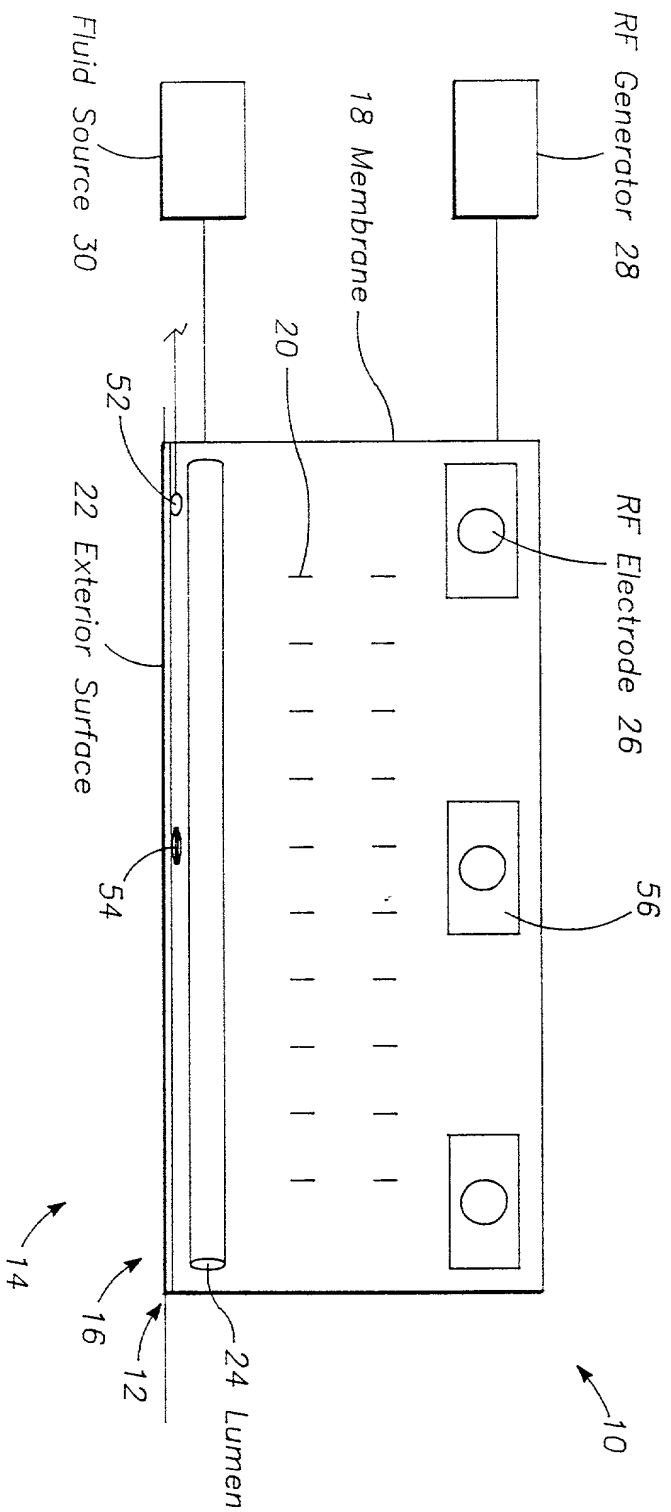


FIG. -1

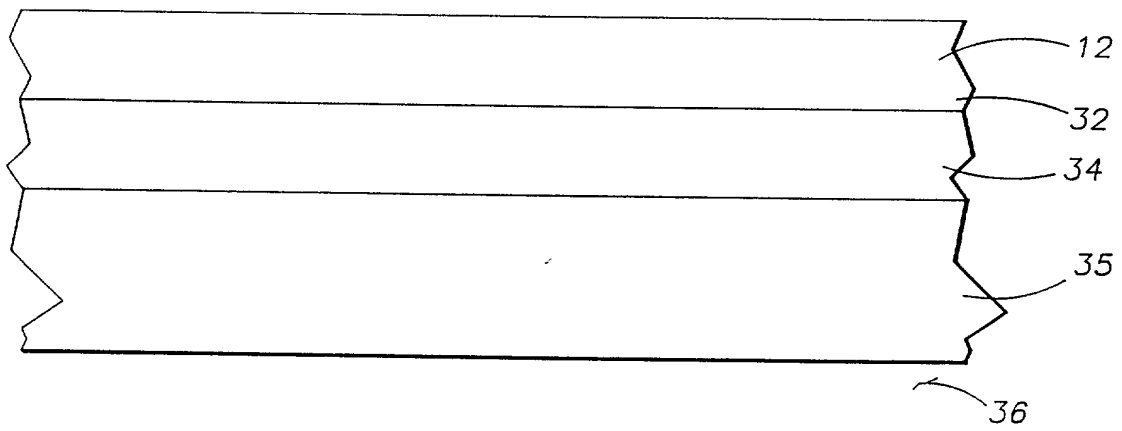


FIG. - 2

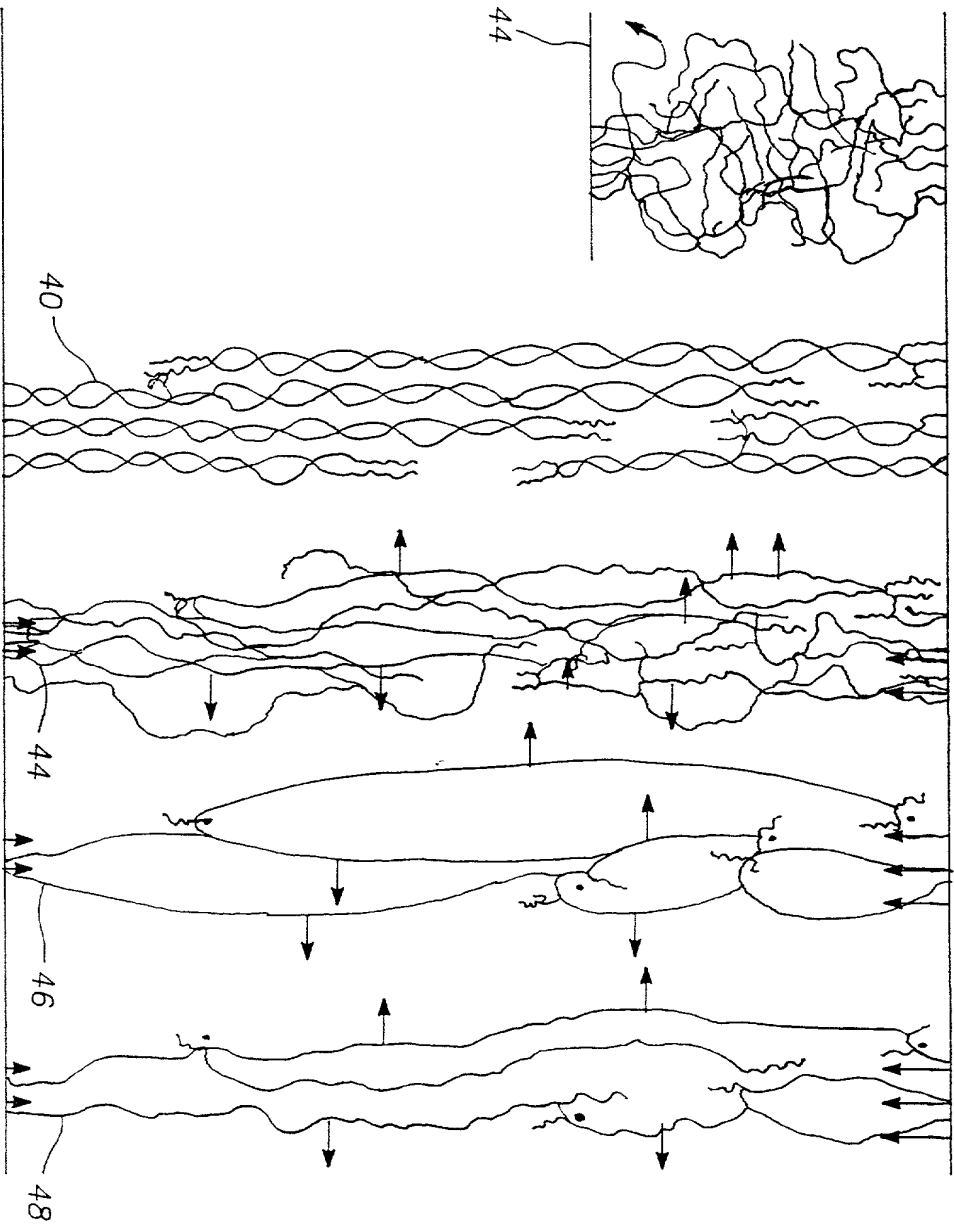


FIG.-3

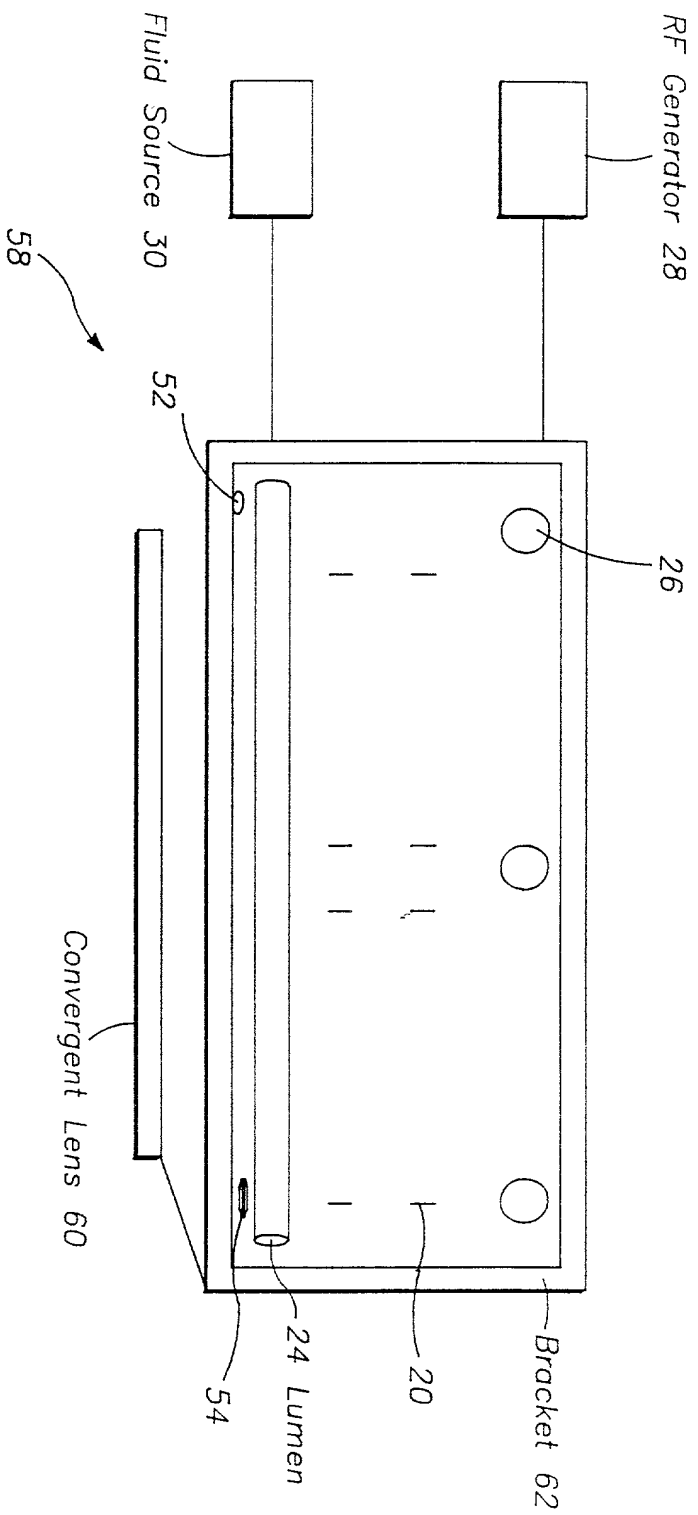


FIG.—4

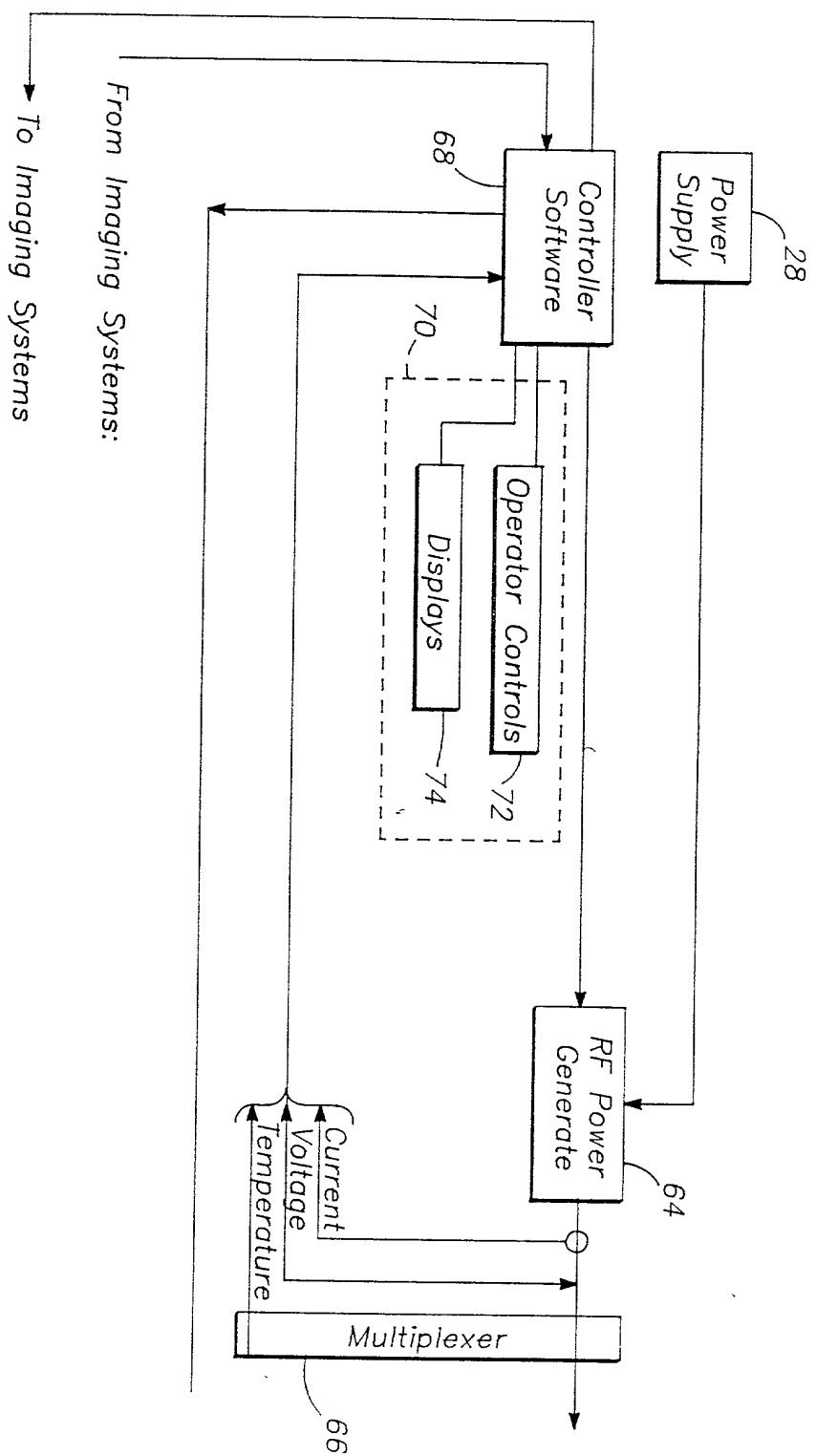


FIG. 5

